Discontinuation of Treatment or Study (Adult)



Patient ID _____ - ___ ID ___ - ____ Date Form Completed: DFCDATE

Instruction: Complete this form when the patient prematurely discontinues study medication, study participation or both.

- 1. Time period: 1 Treatment 2 Follow-up TIMEP
- 2. Is this a discontinuation in study medication and/or study participation? (check all that apply)

□ Study medication (complete Section I) SMED

□ Study participation (complete Section II) **SPART**

SECTION I: STUDY MEDICATION

1. Reason(s) for discontinuing study medication(s) (check all that apply):

	Withdrawal of informed consent RMWCONS	Hypersensitivity reaction RMHYPS
	□ Neutropenia RMNEUT	Pulmonary function impairment RMPF
	Hepatic decompensation RMHDC	Anemia RMANEM
	Autoimmune hepatitis RMAUTO	Renal function impairment RMRF
	Pregnancy RMPREG	Ophthalmologic disorder RMOPH
	Psoriatic lesion RMPSOR	Grade IV toxicity RMTOX4
	Hypoglycemia, hyperglycemia or diabetes mellitus RMDIAB	□ Virological non-response RMVNRSP
	Thyroid disorder/dysfunction RMTHYD	□ Virological breakthrough RMVBRK
	Depression or other psychiatric or mood disorder RMPSY	
	□ Adverse event other than those listed RMAE , specify	RMAES
□ Investigator discretion RMINV, explainRMINVS		
2.	Date of last dose of entecavir (mm/dd/yy): LDEM / LD	ED / LDEY
3.	Date of last dose of peginterferon (mm/dd/yy): LDPM / LD	PD / LDPY

SECTION II: STUDY PARTICIPATION

- 1. Reason(s) for discontinuing study participation *(check all that apply)*:
 - □ Patient lost to follow-up **RSLFUP**
 - □ Withdrawal of informed consent **RSWCONS**
 - □ Patient on alternate therapy for HBV **RSHBVTX**
 - □ Investigator discretion RSINV, explain ______RSINVS_____
 - Other RSOTH, specify _____RSOTHS_____
- 2. Date withdrawn (or considered to be withdrawn) (mm/dd/yy): WDM / WDD / WDY
- 3. Date of last contact (mm/dd/yy): LCM / LCD / LCY